AMENDMENTS TO THE CLAIMS

1. (Currently amended) Pharmaceutical composition for the treatment of acute

and/or chronic pain comprising calcium channel blockers which that are capable of blocking

voltage-dependent calcium channels.

2. (Original) Pharmaceutical composition as defined in claim 1 wherein the

calcium channel is a T-type or L-type channel.

3. (Currently amended) Pharmaceutical composition as defined in claim 1 or 2

for the treatment of allodynia or hyperalgesia.

4. (Currently amended) Pharmaceutical composition according to any one of

claims 1 to 3 claim 1 wherein the calcium channel blocker is mibefradil, its pharmaceutically

acceptable analogues, salts or esters or a dihydropyridine.

5. (Currently amended) Pharmaceutical composition according to claim 1 for the

treatment of pain associated with rheumatoid arthritis, cancer, injuries, back pain, herpes

zoster and post-operative pain.

(Currently amended) Pharmaceutical composition according to any one of 6.

elaims 1 to 5 claim 1 for the topical, oral, parenteral, inhalative or intranasal administration.

(Original) Pharmaceutical composition according to claim 6 in form of an 7.

ointment, gel, crème or a solution or suspension, or plaster.

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8. (Original) Pharmaceutical composition according to claim 6 in form of a nasal

spray or inhalator.

9. (Currently amended) Method for the systemic therapy of pain, comprising

administering a Pharmaceutical pharmaceutical composition according to any one of claims 1

to 3 for the systemic therapy of pain comprising calcium channel blockers that are capable of

blocking voltage-dependent calcium channels.

10. (Currently amended) Pharmaceutical composition according to any one of

elaims 1 to 3 characterised in that claim 1 wherein the composition is in a drug form for oral

administration, wherein the form is selected from the group consisting of a tablet, a capsule, a

coated tablet, a granulate, a juice, a syrup, a suspension, and a solution. tables, capsules,

coated tablets, granulates, juice, syrup, suspensions or solutions are used as oral forms of

application.

11. (Currently amended) Pharmaceutical composition according to any one of

elaims 1 to 3 characterised in that claim 1 wherein the drug form used is formed of comprises

biologically utilizable or biodegradable substances wherein the biological materials are

selected from the group consisting of proteins or proteides, lipids or lipoids, carbohydrates, or

polysaccharides or mixtures of several of such materials and mixtures thereof.

12. (Currently amended) Pharmaceutical composition according to any one of

elaims 1 to 3 characterised in that additionally claim 1 which further comprises one other

pain killer is used.

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13. (Currently amended) Pharmaceutical composition according to claim 12

eharacterised in that wherein the other pain killer used in combination is selected from the

group consisting of an NSAID, a 5HT_{1D} agonist, a dopamin D₂ receptor antagonist, a secale

alcaloid, a beta blocker, a calcium channel blocker-or, and a neurokinin antagonist.

14. (Currently amended) Pharmaceutical composition according to claim 12

characterised in that wherein the NSAID is ibuprofen, meoxicam, indomethacin or naporxen.

15. (Currently amended) Pharmaceutical composition according to claim 12

characterised in that wherein the 5HT_{1D} agonist is sumatriptan, MK-452, naratriptan or 311C.

16. (Currently amended) Pharmaceutical composition according to claim 12

characterised in that wherein the dopamin D₂ receptor antagonist is metoclopramid.

17. (Currently amended) Pharmaceutical composition according to claim 12

characterised in that wherein the secale alcaloid is ergotamin, dihydroergotamin or

metergolin.

18. (Currently amended) Pharmaceutical composition according to claim 12

characterised in that <u>wherein</u> the beta blocker is propranolol or metoprolol.

19. (Currently amended) Pharmaceutical composition according to claim 12

characterised in that wherein the calcium channel blocker is flunarizin or lomerizin.

20. (Currently amended) Pharmaceutical composition according to claim 12

characterised in that wherein the other pain killer to be administered in combination is

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selected from the group consisting of acetylsalicylic acid, paracetamol, clonidin, methysergid, dotarizin, lisurid, pizotifen, valproat, aminotraptilin CP-122,288-or, and UK 116,044.